## Novel electronic states in 5d Ir Oxides produced by strong spin-orbit coupling

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We present a zoology of complex 5d Ir oxides with emphasis on novel interlay of electron correlations, spin-orbit coupling (SOC) and lattice distortion.

5d transition metal compounds are generally believed to be more itinerant than their 3d and 4d analogues. A layered oxide  $Sr_2IrO_4$  with  $Ir^{4+}(5d^5)$ , however, is known to be an antiferromagnetic Mott insulator despite its 4d analogue  $Sr_2RhO_4$  is simply a paramagnetic metal. It was pointed out that the Mott insulating state in this  $Sr_2IrO_4$  is induced by a very strong SOC of ~ 0.5 eV. The large SOC gives rise to a half filled and narrow  $J_{eff}=1/2$  band, leading to a Mott insulting state even in the presence of a weak U~0.5 eV [1]. From a "selection rule" of the magnetic resonant x-ray scattering at Ir L-edge, we unambiguously demonstrated [2] that the ground state of  $Sr_2IrO_4$  is close to a  $J_{eff}=1/2$  antiferromagnet.

Essentially the same  $J_{eff}=1/2$  Mott state was found also in a double layer Ir oxides,  $Sr_3Ir_2O_7A$  number of metallic analogues of  $Sr_2IrO_4$ , including distorted perovskite Sr(Ca) IrO<sub>3</sub> and hexagonal SrIrO<sub>3</sub>, were visited recently. It was unexpectedly found that all compounds are a low carrier density semimetal with an enhanced Wilson ratio  $R_W$ ~10. We argue that those unusual semi-metallic states are formed by an interplay of a large spin orbit coupling and band crossings.

We report also on complex Ir oxides with lattice topology of interest including honeycomb  $Na_2IrO_3$  and spinel  $Ir_2O_4$ , which were theoretically pointed out to be a correlated topological insulator [3] and a Kiteav magnet [4].

[1] B.J.Kim et al., **Phys Rev Lett** 101, 076402 (2008).

[2] B. J. Kim, H. Ohsumi, T. Komesu, S. Sakai, T. Morita, H. Takagi, and T. Arima, Science 323, 1329 (2009).

[3] Shitade et al., Phys. Rev. Lett. 102, 256403 (2009).

[4] J. Chaloupka, G.Jackeli, and G.Khaliullin, Phys. Rev. Lett. 105, 027204 (2010)

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